PATENT SPECIFICATION

NO DRAWINGS

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COMPLETE SPECIFICATION

Mouldable Plastics Compositions based on ethylene copolymer salts

We, FARBENFABRIKEN BAYER AKTIEN-GESELLSCHAFT, a body corporate organised under the laws of Germany, of Leverkusen, Germany, do hereby declare the invention, 5 for which we pray that a patent may be granted to us, and the method by which it is to be performed, to be particularly described in and by the following statement:—

This invention relates to mouldable plas-10 tits compositions comprising carboxylic acids containing more than 5 carbon atoms in admixture with copolymers of ethylene and A-olefinically unsaturated carboxylic

acids containing metal ions.

It is known that copolymers of ethylene and «β-olefinically unsaturated carboxylic acids can be reacted with water-soluble metal compounds. This reaction is accompanied by neutralisation of the carboxyl 20 groups present in the copolymers. In the solid state, plastics compositions of this kind show the same properties as cross-linked polymers although they can be fused, in which state they show the properties of un25 crosslinked polymers. Unfortunately, both

the preparation and the processing of saltlike plastics such as these provide considerable difficulties, particularly when high molecular weight copolymers are used as the 30 starting materials and when compounds of polyvalent metals are employed to neutralise the carboxyl groups present in the copoly-

Such polyvalent metals, for example alka-35 line earth metals, zinc, aluminium or lead, tend to oversolidify the copolymer when used in appreciable quantities to neutralise the carboxyl groups present in it. An excessively intensive "salt crosslinking" occurs, so that 40 the products cannot be processed or can only

be processed at undesirably high tempera-

If salts of the alkali metals are used to neutralise the carboxyl groups present in 45 ethylene copolymers, the difficulties involved in processing are not on the same scale, although the alkali metal carboxylate groups present in the polymers make them undesir-

tures. In some cases, the materials are also

ably over-sensitive to water.

hard and brittle.

It is known from Belgian Patent Specification No. 611,627 that homopolymers and copolymers of olefinically unsaturated monomers can be mixed with 0.05 to 5% by weight, based on the total mixture of car- 55 boxylic acids containing at least four carbon atoms with a view to improving their properties, such as their impact strength and modulus of clasticity.

It is also known from Belgian Patent 60 Specification No. 616,888 that between 0.1 and 5% by weight of carboxylic acids containing 6 to 60 carbon atoms can be added to homopolymers of monoolefins, of ethylene in particular, with a view to improving their 65

Japanese Patent Application No. 16938/ 1961 (published in Chemisches Zentralblatt, 1965, Vol. 21, No. 2808) relates to a mixture of polyethylene with 0.003 to 5% by weight 70 of lithium soaps of fatty acids containing at least 6 carbon atoms, or naphthenic acids. This is said to inhibit crack growth when the polymer is heated for prolonged periods.

By contrast, copolymers of ethylene with 75 2,β-olefinically unsaturated carboxylic acids and optionally other monomers, in which the hydrogen atoms of the carboxyl groups incorporated are completely or partly replaced by monovalent or polyvalent metal 80 ions, are used in the process according to the present invention. The addition of car-

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boxylic acids containing more than 5 carbon atoms, in quantities from 0.1 to 100% by weight based on the copolymer containing metal ions, to an ionically crosslinked and 5 almost unprocessible copolymer of this kind, makes it flexible and eminently processible. It is surprising that homogeneous and clear or transparent plastics compositions showing little or no tendency to disintegrate can be 10 produced. It is assumed that these outstanding properties of the thermoplastic moulding compositions according to the invention are attributable to the interaction between metal ions in the carboxyl-containing copolymer 15 and the carboxyl groups of the carboxylic acids added. The result obtained by the present invention is neither disclosed nor even suggested in the prior art referred to. One object of the present invention is to

20 improve the processing properties of copolymers of ethylene with & B-olefinically saturated carboxylic acids both during and after neutralisation of the carboxyl groups with compounds of polyvalent metals, and in 25 particular the processing of high molecular weight polymers. Another object of the invention is to produce copolymers of ethylene with α,β-tolefinicaly unsaturated carboxylic acids, which have the lowest possible sensi-30 tivity to water. In addition, it is desirable to include in the plastics compositions the inexpensive fatty acids which are available

in large amounts.

Accordingly, the present invention relates 35 to a mouldable plastics composition comprising a copolymer of ethylene and an a, ßolefinically unsaturated carboxylic acid and optionally other comonomers, from 10 to 100% of the hydrogen atoms in the carboxyl 40 groups of the copolymer being replaced by monovalent or polyvalent metal ions, whilst the copolymers free of metal ions has a melt index of at least 0.2g/10 mins., which composition contains from 0.1 to 100% by weight

45 based on the copolymer containing metal ions, of a carboxylic acid containing more than 5 carbon atoms.

It is surprising that homogeneous and clear or transparent plastics compositions 50 showing no tendency to disintegrate can be obtained by combining the aforementioned salt-like copolymers with carboxylic acids dontaining more than 5 carbon atoms because in general, carboxylic acids of this 55 kind are not completely compatible with the aforementioned copolymers. It is also surprising that the addition of carboxylic acids containing more than 5 carbon atoms, in particular aliphatic carboxylic acids con-60 taining more than 10 carbon atoms (for example, oleic acid) to hard, ionically cross-

linked and almost unprocessible polymeric salts of polyvalent metals (for example the calcium salt of a high molecular weight 65 ethylene-acrylic acid copolymer) makes the

polymeric salts flexible and eminently processible. Neither was it expected that a polymeric salt of this kind, which may be regarded as being completely insoluble both in aqueous and in most organic solvents, 70 would readily absorb the fatty acid added, which becomes integrated as a constituent of the polymer. This is apparent inter alia from the fact that the plastics compositions obtained are completely clean, whilst the 75 carboxylic acid incorporated shows no tendency to exude, except when added in large quantities, and even then its tendency to exude is still very limited. Finally, it has also been found that the addition of rela- 80 tively large quantities of carboxylic acids containing more than 5 carbon atoms is accompanied by little or no deterioration in the ultimate tensile strength of the salt-like polymers, whilst on the other hand their 85 breaking elongation is favourably increased. Ethylene copolymers containing at least

40 mol % and preferably 50 to 96 mol % of copolymerised ethylene; 1 to 30 mol % and preferably 2 to 20 mol % of copiolymerised carboxylic acid and, optionally, and the state of the state polymerised units of other monomers, can advantageously be used as the copolymers of ethylene with a β -olefinically unsaturated carboxylic acids. The melt index of the co- 95 polymers should be at least 0.2 g/10 mins and may even exceed 100 g/10 mins. The melt index is determined in accordance with

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In addition to the a, \beta-olefinically unsatur- 100 ated carboxylic acid(s) and ethylene, the ethylene copolymers of the aforementioned type may also contain further comonomers, in proportions up to 30 mol %.

The following are examples of such co- 105

a) Derivatives of α,β-olefinically unsaturated monocarboxylic acids, such as acrylamide, methacrylamide, acrylonitrile, methacrylonitrile and esters of acrylic and methacrylic 110 acid, particularly those with saturated monohydric aliphatic alcohols containing 1 to 20 carbon atoms cycloaliphatic alcohols containing 5 to 7 carbon atoms, phenols, aralkyl alcohols or heterodyclic alcohols, such as 115 esters of the aforementioned acids with methyl, ethyl, propyl, isopropyl, isobutyl, hexyl, octyl or stearyl alcohol, cyclohexanol, methyl-cyclohexanol, benzyl alcohol, phenol, cresol or furfuryl alcohol.

Monoesters of αβ-mono-olefinically unsaturated monocarboxylic acids containing 3 to 4 carbon atoms with dihydric saturated aliphatic alcohols containing 2 to 4 carbon atoms, for example 2-hydroxyethyl meth- 125 acrylate, 2-hydroxypropyl methacrylate, 4hydroxy-n-butyl methacrylate, 2-hydroxyethl acrylate, 2-hydroxypropyl acrylate and 4-hydroxy-n-butyl acrylate, can also be em-

ployed.

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b) aromatic vinyl and vinylidene compounds, such as styrene, a-methylstyrene, vinyl toluene, p-chlorostyrene or other nuclear-substituted vinyl benzenes;

5 c) Esters of vinyl alcohol with carboxylic acids containing 2 to 18 carbon atoms or with hydrohalic acids, vinyl ethers and vinyl ketones, such as vinyl chloride, vinylidene chloride, vinyl acetate, vinyl propionate, in vinyl stearate, vinyl ethyl ether, vinyl isobutyl ether, or vinyl amides such as vinyl formaldehide and vinyl acetamide:

d) Conjugated dienes containing 4 to 6 carbon atoms such as butadiene, isoprene, 2,3-

15 dimethyl butadiene and chloroprene; and e) Propylene, cycloheptatriene or norborna-

Acrylic esters, methacrylic esters and the vinyl esters listed are preferably used as the 20 additional comonomers.

Although ethylene copolymers incorporating these additional monomers may be used, it is preferred to use copolymers of ethylene with 2β-olefinically unsaturated carboxylic

25 acids only.

The elefinically unsaturated carboxylic acids can be incorporated in the polymer in quantities from 1 to 30 mol % and preferably in quantities from 2 to 20 mol %. Pre-30 ferred carboylic acids of this kind include acrylic acid and methacrylic acid, although it is also possible to use itaconic acid, maleic acid of fumaric acid or their half esters with alcohols containing 1 to 8 carbon atoms, for 35 example monoethyl aleate or fumarate.

It is also possible to use, for the plastics compositions according to the invention, copolymers of ethylene in which the carboxyl group has been introduced by a subsequent 40 operation, for example, oxidation, grafting, hydrolysis or saponification, or by the reaction of an organometallic derivative of the polymer with CO2 or by the reaction of a reactive group to form the carboxyl groups 45 (for example the reaction of an OH-group with an acid anhydride of a polyvalent carboxylic acid (to form the half ester.).

The copolymers of ethylene with z, \betaolefinically unsaturated carboxylic acids and 50 optionally further comonomers, can be prepared by methods known per se, for example at pressures from 50 to 300 atms. and at temperatures in the range from 50 to 300°C, by ionic or free radical polymerisation, as de-55 scribed for example in Belgium Patent Specification 679,490, in German Patent Specification Nos. 650,038, 745,425 or in British Patent Specification No. 963,380.

Preparation of the metal salts by neutral-60 ising the above-described copolymers of ethylene and a B-olefinically unsaturated carboxylic acids and optionally other monomers, may be carried out under the conditions of the process disclosed in British

65 Patent Specification No. 1,011,981.

The carboxyl groups of the copolymers are either completely or partly neutralised although at least 10% of the carboxyl groups present are neutralised.

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Suitable neutralising agents include the 70 water-soluble or difficultly water-soluble compounds of mono- and/or polyvalent metals of Groups I, II, IV-A and VIII of the Periodic System, mentioned in British Patent Specification No. 1,011,981 (cf. Hand. 75 book of Chemistry and Physics, Chemical Rubber Publishing Company, 37th Edition, page 392).

Suitable carboxylic acids containing more than 5 carbon atoms for addition to the polymeric salts include saturated or unsaturated aliphatic, cycloaliphatic, araliphatic, aromatic and heterocyclic carboxylic acids or carboxylic acid mixtures of natural or synthetic origin with molecular weights in excess 85 of 103, for example naphthenic acids, synthetic fatty acids, cottonseed oil acids, tall oil fatty acids, abietic acid, resin acids, cinnamic acid, phenylbutyric acid, phenylacetic acid, salicyclic acid, ω-aminocaproic acid, 90 phnylalanine or N-cyclohexylaminosuccinic acid; it is preferred, however, to use saturated or unsaturated aliphatic monocarboxylic or polycarboxylic acids.

The following are examples of suitable 95 saturated aliphatic fatty acids: caprylic acid, lauric acid, myristic acid, arachic acid, behenic acid, cerotic acid, palmitic acid and

stearic acid

In addition, oleic acid, decylenic acid, 100 palymitoleic acid, dodecylenic acid, linolic acid, linoleic acid, ricinoleic acid, eleostearic acid, parinaric acid, arachidonic acid, cetoleic acid and nervonic acid, are examples of suitable unsaturated aliphatic fatty acids.

Examples of aliphatic polycarboxylic acids are adipic acid, pimelic acid, suberic acid, azelaic acid, undecane dicarboxylic or higher polycarboxylic acids obtained by polymerisation or oxidation processes or, for 110 example, by the copolyerisation of unsaturated carboxylic acids with olefins, vinyl or acrylic compounds or from petrochemicals or by the oligomerisation of natural or synthetic fatty acids, for example so-called di- 115 meric acids.

Carboxylic acids or polycarboxylic acids obtained by polyaddition or polycondensation are also suitable, for example saturated and unsaturated acid polyesters, polyethers, 120 polyamnies or polyurethanes containing carboxyl groups.

The carboxylic acids containing more than 5 carbon atoms are added to the salt form of the ethylene copolymers in quantities from 125 0.1 to 100% by weight, based on polymer containing metal ions, although they are preferably added in quantities from 3 to 50% by weight. Aliphatic carboxylic acids containing more than 10 carbon atoms are per-

ferably used. Incorporation may be effected by combining the polymer salt and the carboxylic 5 acid in solution, in emulsion or in the solid state at temperatures in the range from 10 to 280°C, for example by directly working the carboxylic acid in dissolved, emulsified or fused form into a dispersion or emulsion 10 of the polymer salt. It is preferred, however, to work the carboxylic acid into the salt-like polymer on mixing rolls or in a kneader at temperatures above 50°C. An alternative method of incorporating the car-15 boxylic acid comprises adding it during preparation of the metal salt of the ethylene co-This alternative eliminates one stage and simplifies handling of the plastics compositions, because the required material 20 is directly formed.

The products obtained by the process according to the invention may be used either in bulk, in solution or in emulsion, for example, for coating purposes. As solids they 25 are suitable as elastomers, extrusion or injection-moulding compositions. Highly transparent coating films and mouldings with a high resistance to chemicals, coupled with a high resistance to chemical and processing pro-favourable mechanical and processing pro-30 perties, can be prepared from them.

products can be stretched or spun.

Example 1: The 100% neutralised sodium salt of a copolymer of ethylene and 10% by weight 35 of polymerised methacrylic acid, prepared in accordance with British Patent Specification No. 1,011,981, page 10, (product No. 2 from Table IV), is used: it does not show any tendency to flow in a melt-index apparatus. 40 The material is difficult or totally impossible to process in extruders or injection-moulding machines. These difficulties are caused by the extremely low melt index of the polymer salt. It is intended in the following to de-45 monstrate how the melt index of the material is improved by the addition of stearic acid, thus eliminating any difficulties in its pro-For this purpose, stearic acid is worked in various quantities into separate 50 portions comprising 100 parts by weight of the polymer salt on a grooved roller at 120°C, and the improvement in processibility is followed by reference to the melt index. In every instance, transparent compositions

55 are obtained which can be processed into injection mouldings or films. Equivalent of acid Melt index % by weight per equivalent of stearic acid Na in the polymer 0.0 g/10 mins.

| 0 | 0.0 - 5.5 11 22 | approx. 0.15 0.33 0.33 | 0.8 2.2 5.5 6.5 |
|----|--------------------------|------------------------------|--------------------------|
| 55 | 33 50 | ., 1.0 ., 1.5 | 6.5 |

Example 2:

The corresponding neutral calcium salt is used in place of the sodium salt of the polymer employed in Example 1. The composition, which does not show any flow in the 70 melt-index apparatus, is difficult to fuse and cannot be processed. Oleic acid is worked in various quantities into this material on a toothed roll whose surface temperature is

above 120°C. Equivalent of oleic % by weight acid per equivalent Melt index of oleic acid of Ca in the polymer g/10 mins.

| CIC acid | 0.0 | no ilow | |
|----------|------|------------|---|
| 0.0 | | 0.3 | Ì |
| 5.9 | 0.16 | 1.2 | |
| 11 | 0.3 | | |
| | 0.65 | 4.0 | |
| 24 | 1.00 | 5.0 | |
| 36.7 | 1.00 | can now be | |

The transparent material can now be processed into films and mouldings.

Example 3: The 100% neutralised zinc salt of a copolymer of ethylene and 3.8% by weight of maleic acid is prepared by rolling the copolymer with zinc acetate at 120°C, followed 90 by removal of the acetic acid liberated in a vacuum dryer at 150°C. This is a material which cannot be processed and which does not flow in the melt index apparatus. 100 parts by weight thereof are kneaded with 26 95 parts by weight of ricinoleic acid for 30 min-utes at 140°C in a nitrogen atmosphere, resulting in the formation of a clear, satisfactorily processible polymer composition with a melt index of 15 g/10 mins. 100

Metal salts of the following types of poly-Example 4: mer are used in this and the following Ex-

Type 1: Copolymer of ethylene and acrylic 105 acid with a polymerised acrylic acid content of 19.5% by weight and a melt index of 120

Type 2: Copolymer of ethylene and methacrylic acid with a polymerised methacrylic 110 acid content of 31% by weight and a mult

index of 18 g/10 mins. Type 3: Copolymer of 9.5% by weight of acrylic acid, 10% by weight of ethyl acrylate and 80.5% by weight of ethylene, with 115 a melt index of 15 g/10 mins.

The 100% neutralised calcium salts of polymers 1 and 2 do not flow in the melt index apparatus. After 15% by weight of oleic acid have been worked in on a kneader 120 at 120°C, the melt indices of these polymers are 9.5 and 1.8 g/10 mins, respectively. The clear polymer compositions can now be processed.

Example 5: The 65% neutralised aluminium salt of polymer 3 (Example 4) is a highly crosslinked, infusible mass. After 15% by weight of linoleic acid (approx. 0.5% equivalent of acid per equivalent of aluminium 130

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in the polymer) have been worked into it, the polymer salt loses its crosslinked character, is transparent, satisfactorily processible and has a melt index of 2.8 g/10 mins.

5 Example 6:

The effect of various types of carboxylic acids is demonstrated in the following, with reference to the 100% neutralised calcium salt of polymer 1 (Example 4). A kneader 10 was used for working in which was carried out at 150°C. in the absence of air:—

| 0/1 | Melt index |
|-----------------------------|------------|
| % by weight carboxylic acid | g/10 min: |
| 0.0 | no flow |
| 15 10 stearie acid | 11 |
| 5.0 adipic acid | 9 |
| 8.0 azelaic acid | 14 |
| 15.0 abietic acid | 2.9 |
| 9.0 phenylacetic acid | 16 |
| 20 11 e-aminocaproic acid | 8 |
| 8 copolymer of 37 mol | Ū |
| % of acrylic acid and | |
| and 63 mol % of ethy- | |
| lene | 1.2 |
| 25 Example 7. | 1.5 |

A polymer of 90% by weight of ethylene and 10% by weight of methacrylic acid, 70% of which is in the form of the sodium salt and which has a breaking elongation of 30 490%, is kneaded with stearic acid at 120°C.

After storage for 5 days, the following breaking elongations are measured.

| % by weight | Breaking |
|-----------------|--------------|
| of stearic acid | elongation % |
| 35 — | 490 |
| .5 | 540 |
| 10 | 551 |

The increase in breaking elongation improves the blowing properties of the material, 40 and the elastic properties of the films produced from it.

WHAT WE CLAIM IS:—

1. A mouldable plastics composition com-

prising a copolymer of ethylene and an z, \(\beta\)olefinically unsaturated carboxylic acid and 45
optionally other comonomers, from 10 to
100% of the hydrogen atoms in the carboxyl groups of the copolymer being replaced by monovalent or polyvalent metal
ions, whilst the copolymer free of metal ions 50
has a melt index of at least 0.2 g/10 mins,
which composition contains from 0.1 to
100% by weight, based on the copolymer
containing metal ions, of a carboxylic acid
containing more than 5 carbon atoms.

2. A mouldable plastics composition as claimed in claim 1, which contains from 3 to 50% by weight, based on the copolymer containing metal ions, of a carboxylic acid containing more than 5 carbon atoms.

3. A mouldable plastics composition as claimed in claim 1 or 2, wherein the carboxylic acid containing more than 5 carbon atoms is an aliphatic carboxylic acid containing more than 10 carbon atoms.

4. A mouldable plastics composition as claimed in any of claims 1 to 3 wherein the copolymer comprises at least 40 mol % of ethylene, 1 to 30 mol % of an α,β-olefinically unsaturated carboxylic acid and optionally 70 up to 30 mol % of other comonomers, the hydrogen atoms in the carboxyl groups of the copolymer being completely or partly replaced by ions of a metal of Group I, II, III, IV-A or VIII of the Periodic System (as 75 hereinbefore specified).

5. A mouldable plastics composition as claimed in claim 1 substantialy as described with reference to any of the Examples.

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